	FILE	'REGISTRY' ENTERED AT 12:35:55 ON 06 OCT 2008
		EXP LANSOPRAZOLE/CN
L1		1 S E3
		EXP ETHYLENEDIAMINETETRAACET/CN
L2		1 S E5
		EXP N-METHYLGLUC/CN
L3		1 S E4
	FILE	'HCAPLUS' ENTERED AT 12:37:02 ON 06 OCT 2008
L4		0 S L1 AND L2
		IDECTORDAL ENGEDED AT 10 27 02 ON 06 OOT 0000
	FILE	'REGISTRY' ENTERED AT 12:37:23 ON 06 OCT 2008
L5		EXP ETHYLENEDIAMINETETRAACET/CN 1 S E10
ьэ		EXP ETHYENEDIAMINETETRAACETIC ACID SOD/CN
		EXP ETHYLENEDIAMINETETRAACETIC ACID SOD/CN
L6		1 S E4
по		1 3 4
	FILE	'HCAPLUS' ENTERED AT 12:38:45 ON 06 OCT 2008
L7		0 S L1 AND L6
L8		15 S L1 AND L5
L9		7 S L8 AND (PY<2004 OR AY<2004 OR PRY<2004)
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=> file registry
COST IN U.S. DOLLARS
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FULL ESTIMATED COST

E12

1

SINCE FILE TOTAL
ENTRY SESSION
0.21 0.21

FILE 'REGISTRY' ENTERED AT 12:35:55 ON 06 OCT 2008
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

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STRUCTURE FILE UPDATES: 5 OCT 2008 HIGHEST RN 1057399-47-9 DICTIONARY FILE UPDATES: 5 OCT 2008 HIGHEST RN 1057399-47-9
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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

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=> exp lansoprazole/cn
E1
                   1 LANSOPEP/CN
E.2
                    1
                             LANSOPHED/CN
                   1 --> LANSOPRAZOLE/CN
E3
                 LANSOPRAZOLE/CN
LANSOPRAZOLE CALCIUM/CN
LANSOPRAZOLE SODIUM/CN
LANSOPRAZOLE SULFIDE/CN
LANSOPRAZOLE SULFONE/CN
LANSOPRAZOLE-AMOXICILLIN MIXT./CN
LANSOPRAZOLE-CLARITHROMYCIN MIXT./CN
LANSOPRAZOLE-LEVOFLOXACIN MIXT./CN
LANSOPRAZOLE-SITAFLOXACIN MIXT./CN
LANSOPRID/CN
E4
E5
Ε6
E7
E8
E9
E10
E11
E12
=> s e3
                    1 LANSOPRAZOLE/CN
T.1
=> exp ethylenediaminetetraacet/cn
                              ETHYLENEDIAMINETETRA (METHYLPHOSPHONIC ACID) MONOHYDRATE/CN
E1
                    1
E2
                     1
                              ETHYLENEDIAMINETETRA-3-PROPIONIC ACID/CN
Е3
                     0 --> ETHYLENEDIAMINETETRAACET/CN
E4
                    1
                            ETHYLENEDIAMINETETRAACETAMIDE/CN
E5
                     1
                              ETHYLENEDIAMINETETRAACETATE/CN
                            ETHYLENEDIAMINETETRAACETATOCOBALTATE(III)/CN
E6
                     1
                           ETHYLENEDIAMINETETRAACETATOINDATE(1-)/CN
ETHYLENEDIAMINETETRAACETATOVANADATE(1-)/CN
ETHYLENEDIAMINETETRAACETDIDODECYLIMIDE/CN
ETHYLENEDIAMINETETRAACETIC ACID/CN
ETHYLENEDIAMINETETRAACETIC ACID ANHYDRIDE-ETHYLENEDIAMINE CO
E7
                     1
E8
                    1
E9
                    1
E10
                    1
E11
                   1
                             POLYMER/CN
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ETHYLENEDIAMINETETRAACETIC ACID BARIUM SALT/CN

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1 ETHYLENEDIAMINETETRAACETATE/CN
T.2
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                          N-METHYLGLAUCINE IODIDE/CN
                  1
E2
                  1
                          N-METHYLGLAUDINIUM IODIDE/CN
E3
                   0 --> N-METHYLGLUC/CN
E4
                  1
                         N-METHYLGLUCAMINE/CN
E5
                 1
                         N-METHYLGLUCAMINE 5,5'-(DODECANEDIOYLDIIMINO)-BIS(2,4,6-TRII
                         ODO-N-METHYLISOPHTHALAMATE)/CN
                  1
                         N-METHYLGLUCAMINE ACETYLSALICYLATE/CN
                1 N-METHYLGLUCAMINE ACEITESABICIEMIE, C.
1 N-METHYLGLUCAMINE ANTIMONATE/CN
1 N-METHYLGLUCAMINE CYCLOHEXYLSULFAMATE/CN
1 N-METHYLGLUCAMINE DIATRIZOATE/CN
1 N-METHYLGLUCAMINE HYDROCHLORIDE/CN
1 N-METHYLGLUCAMINE IODOMETHANESULFONATE/CN
1 N-METHYLGLUCAMINE IOTHALAMATE/CN
E7
E8
E9
E10
E11
E12
=> s e4
L3
                   1 N-METHYLGLUCAMINE/CN
=> file hcaplus
COST IN U.S. DOLLARS
                                                                       SINCE FILE
                                                                                             TOTAL
                                                                              ENTRY
                                                                                         SESSION
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FILE 'HCAPLUS' ENTERED AT 12:37:02 ON 06 OCT 2008
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FILE COVERS 1907 - 6 Oct 2008 VOL 149 ISS 15 FILE LAST UPDATED: 5 Oct 2008 (20081005/ED)

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

```
=> s 11 and 12
1904 L1
100 L2
L4 0 L1 AND L2
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=> file registry
COST IN U.S. DOLLARS

FULL ESTIMATED COST

=> s e5

SINCE FILE TOTAL

16.37

16.58

FULL ESTIMATED COST ENTRY SESSION 2.69 19.27

FILE 'REGISTRY' ENTERED AT 12:37:23 ON 06 OCT 2008 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2008 American Chemical Society (ACS)

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STRUCTURE FILE UPDATES: 5 OCT 2008 HIGHEST RN 1057399-47-9 DICTIONARY FILE UPDATES: 5 OCT 2008 HIGHEST RN 1057399-47-9

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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=> avn athul	anadis	minetetraacet/cn
E1		ETHYLENEDIAMINETETRA (METHYLPHOSPHONIC ACID) MONOHYDRATE/CN
E2	1	ETHYLENEDIAMINETETRA-3-PROPIONIC ACID/CN
E3	_	ETHYLENEDIAMINETETRAACET/CN
E4	1	ETHYLENEDIAMINETETRAACETAMIDE/CN
E5	1	ETHYLENEDIAMINETETRAACETATE/CN
E6	1	ETHYLENEDIAMINETETRAACETATOCOBALTATE(III)/CN
E7	1	ETHYLENEDIAMINETETRAACETATOINDATE(1-)/CN
E8	1	ETHYLENEDIAMINETETRAACETATOVANADATE(1-)/CN
E9	1	ETHYLENEDIAMINETETRAACETDIDODECYLIMIDE/CN
E10	1	ETHYLENEDIAMINETETRAACETIC ACID/CN
E11	1	ETHYLENEDIAMINETETRAACETIC ACID ANHYDRIDE-ETHYLENEDIAMINE CO
		POLYMER/CN
E12	1	ETHYLENEDIAMINETETRAACETIC ACID BARIUM SALT/CN
=> s e10		
L5	1 "ET	HYLENEDIAMINETETRAACETIC ACID"/CN
		ninetetraacetic acid sod/cn
E1	1	ETHYDINE/CN
E2	1	ETHYDRONATE/CN
<b>E</b> 3	-	ETHYENEDIAMINETETRAACETIC ACID SOD/CN
<b>E</b> 4	1	ETHYL/CN
<b>E</b> 5	1	ETHYL (((((1S)-1-(1H-BENZIMIDAZOL-2-YL)-2-(4-(1,1-DIOXIDO-3-
		OXOISOTHIAZOLIDIN-5-YL) PHENYL) ETHYL) AMINO) CARBONYL) AMINO) ACE
		TATE TRIFLUOROACETATE/CN
<b>E</b> 6	1	ETHYL ((((2-((4-CHLOROPHENYL)OXY)ETHYL)AMINO)CARBONYL)OXY)AC
		ETATE/CN
<b>E</b> 7	1	ETHYL ((((2-(3-PHENYLOXYPHENYL)ETHYL)AMINO)CARBONYL)OXY)ACET
		ATE/CN
E8	1	ETHYL ((((5-(((TERT-BUTOXYCARBONYL)AMINO)METHYL)-6-ISOBUTYL-
		2-METHYL-4-(4-METHYLPHENYL)PYRIDIN-3-YL)AMINO)CARBONYL)OXY)A
		CETATE/CN

E9	1	ETHYL ((((5-(AMINOMETHYL)-6-ISOBUTYL-2-METHYL-4-(4-METHYLPHE NYL)PYRIDIN-3-YL)AMINO)CARBONYL)OXY)ACETATE DIHYDROCHLORIDE/CN
E10	1	ETHYL ((((TERT-BUTOXYCARBONYL)AMINO)SULFONYL)(4-((2S)-2-((TERT-BUTOXYCARBONYL)AMINO)-2-(5-(TRIFLUOROMETHYL)-1H-BENZIMIDAZOL-2-YL)ETHYL)-2-CHLOROPHENYL)AMINO)ACETATETRIFLUOROACETATE/CN
E11	1	ETHYL ((((TERT-BUTOXYCARBONYL)AMINO)SULFONYL)(4-((2S)-2-((TERT-BUTOXYCARBONYL)AMINO)-2-(5-(TRIFLUOROMETHYL)-1H-BENZIMIDAZOL-2-YL)ETHYL)PHENYL)AMINO)ACETATE TRIFLUOROACETATE/CN
E12	1	ETHYL (((1-((CYCLOHEXYLAMINO)CARBONYL)PIPERIDIN-4-YL)METHYL) (4-((DODECYLAMINO)CARBONYL)BENZYL)AMINO)(OXO)ACETATE/CN
=> exp ethvl	onedia	minetetraacetic acid sod/cn
E1	1	ETHYLENEDIAMINETETRAACETIC ACID N, N'-DIOXIDE/CN
E2	1	ETHYLENEDIAMINETETRAACETIC ACID POTASSIUM SALT/CN
E3	0>	ETHYLENEDIAMINETETRAACETIC ACID SOD/CN
E4	1	ETHYLENEDIAMINETETRAACETIC ACID SODIUM SALT/CN
E5	1	ETHYLENEDIAMINETETRAACETIC ACID SYM-BIS (N-(B-MERCAPTOET
		HYL)AMIDE)/CN
E6	1	ETHYLENEDIAMINETETRAACETIC ACID SYM-BIS(N-(B-MERCAPTOET HYL)AMIDE), POLYMER/CN
E7	1	ETHYLENEDIAMINETETRAACETIC ACID SYM-BIS(N-METHYLAMIDE)/CN
E8	1	ETHYLENEDIAMINETETRAACETIC ACID TETRAAMMONIUM SALT/CN
E9	1	ETHYLENEDIAMINETETRAACETIC ACID TETRAHYDRAZIDE/CN
E10	1	ETHYLENEDIAMINETETRAACETIC ACID TETRAPOTASSIUM SALT/CN
E11	1	ETHYLENEDIAMINETETRAACETIC ACID TETRASODIUM SALT/CN
E12	1	ETHYLENEDIAMINETETRAACETIC ACID TRIAMMONIUM SALT/CN
=> s e4		
L6	1 "ET	HYLENEDIAMINETETRAACETIC ACID SODIUM SALT"/CN
=> file hcap		
COST IN U.S.	DOLLA	
		ENTRY SESSION

11.22

30.49

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FILE COVERS 1907 - 6 Oct 2008 VOL 149 ISS 15 FILE LAST UPDATED: 5 Oct 2008 (20081005/ED)

FULL ESTIMATED COST

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

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=> s 11 and 16
           1904 L1
           1335 L6
L7
               0 L1 AND L6
=> s 11 and 15
           1904 L1
          34216 L5
             15 L1 AND L5
1.8
=> s 18 and (PY<2004 or AY<2004 or PRY<2004)
       24009717 PY<2004
        4787037 AY<2004
        4258419 PRY<2004
L9
               7 L8 AND (PY<2004 OR AY<2004 OR PRY<2004)
=> d 19 1-7 ti abs bib
     ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
ΤI
     Injectable composition comprising lansoprazole and a chelating agent
AΒ
     An injectable composition comprising a combination of lansoprazole, its
     optically active isomer, or a salt thereof, and a chelating agent, which
     is used at pH 9 to 12. The injectable composition has excellent stability and
     solubility, and has such a high-quality that particulate insolubles are not
     formed when the composition is kept and supplied in a glass container and even
     in a plastic container and also when the composition is kept in these
     containers for a long time. Thus, a composition contained lansoprazole 30,
     N-methylglucamine 10, mannitol 60, and NaOH 3.45 mg and water for
     injection 5 mL.
     2005:567110 HCAPLUS <<LOGINID::20081006>>
ΑN
DN
     143:65527
ΤI
     Injectable composition comprising lansoprazole and a chelating agent
ΙN
     Doen, Takayuki; Inoue, Tomoko
PA
     Takeda Pharmaceutical Company Limited, Japan
SO
     PCT Int. Appl., 58 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 1
     PATENT NO.
                           KIND
                                     DATE APPLICATION NO. DATE
     WO 2005058277
                            A1 20050630 WO 2004-JP18956 20041213 <--
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               CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
               GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
               LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
          NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MD, NE, SN, TD, TC
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IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS

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US 20070191286 A1
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PRAI JP 2003-419288
                               20031217 <--
    WO 2004-JP18956
                        W
                               20041213
             THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 5
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L9
    ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
ΤI
    Film comprising therapeutic agents
AΒ
    The present invention is related to the composition and methods of manufacture
of
    orally-dissolvable, edible films as a vehicle for the non-invasive
    administration of nitroglycerin, as well as other therapeutic agents
    either with or without nitroglycerin, through the mucosal tissues of the
    oral cavity. The films include a water-soluble film-forming polymer, such as
    pullulan. Methods for producing the films are also disclosed.
    2005:55116 HCAPLUS <<LOGINID::20081006>>
ΑN
    142:141267
DN
ΤI
    Film comprising therapeutic agents
    Maibach, Todd
IN
PA
    USA
SO
    PCT Int. Appl., 49 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LA
FAN.CNT 1
                               DATE
                       KIND
                                                                 DATE
    PATENT NO.
                                         APPLICATION NO.
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            GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
            LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
            NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
            TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
        RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
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            SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
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                         A1
PRAI US 2003-484009P
                         Р
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    US 2003-497426P
                        Р
                               20030821 <--
    WO 2004-US21038
                        W
                               20040630
    ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
L9
TI
    Fast dissolving orally consumable films containing a sweetener
    A consumable film adapted to adhere to and dissolve in the oral cavity,
    comprises at least one water-soluble polymer, a taste-masking effective amount
    of a sweetener, and a pharmaceutically active agent having a sufficiently
    unpleasant taste that it is desirably masked by the sweetener. For
    example, a buccal film was formulated containing dextromethorphan · HBr
    22.7322, Amberlite IRP69 24.2477, xanthan gum 0.1165, locust bean gum
    0.1365, carrageenan 0.5851, pullulan 31.2066, K sorbate 0.1170, menthol
    3.908, peppermint flavor 0.3908, cherry flavor 0.3908, sour cherry 3.3871,
    Warm Sensation 0.8362, artificial masking flavor 0.6273, Succulence
```

```
0.3908, FD&C Red Number 40 0.0149, polysorbate 80 0.6826, Atmos 300 0.6826,
     glycerin 2.9256, mannitol 3.9008, and sucralose 2.7279 %.
     2003:892252 HCAPLUS <<LOGINID::20081006>>
ΑN
DN
     139:354513
     Fast dissolving orally consumable films containing a sweetener
TΙ
ΙN
     Kulkarni, Neema; Kumar, Lori D.; Sorq, Albert
PA
SO
     U.S. Pat. Appl. Publ., 14 pp., Cont.-in-part of U.S. Ser. No. 395,104.
     CODEN: USXXCO
DT
     Patent
     English
LA
FAN.CNT 3
     PATENT NO.
                        KIND
                                DATE
                                            APPLICATION NO.
                                                                    DATE
                         ____
     US 20030211136
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PΤ
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     CA 2520986
                                20071113
     CA 2572461
                         A1
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                          A2
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     EP 1676557
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             LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
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- L9 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
- TI Preparation of aqueous clear solution dosage forms with bile acids
- AB Compns. for pharmaceutical and other uses comprise clear aqueous solns. of bile acids which do not form any detectable ppts. over selected ranges of pH values of the aqueous solution The compns. comprise (i) water, (ii) a bile acid component in the form of a bile acid, bile acid salt, or a bile acid conjugated with an amine by an amide linkage; and (iii) either or both an aqueous soluble starch conversion product and an aqueous soluble non-starch polysaccharide. The composition remains in solution without forming a precipitate over a

range of pH values and, according to one embodiment, remains in solution for all pH values obtainable in an aqueous system. The composition may further contain

a pharmaceutical compound, such as insulin, heparin, bismuth compds., amantadine and rimantadine. For example, solution dosage forms that did not show any precipitation at any pH were prepared containing ursodeoxycholic acid (UDCA) 22

g, 1N NaOH 75 mL, chenodeoxycholic acid (CDCA) 3 g, maltodextrin 875 g, bismuth citrate 4 g, citric acid or lactic acid as needed, and purified water to make 1  $\rm L$ .

- AN 2002:185616 HCAPLUS <<LOGINID::20081006>>
- DN 136:252482
- TI Preparation of aqueous clear solution dosage forms with bile acids
- IN Yoo, Seo Hong
- PA USA
- SO U.S. Pat. Appl. Publ., 35 pp., Cont.-in-part of U.S. 6,251,428. CODEN: USXXCO
- DT Patent
- LA English
- FAN.CNT 5

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RE.CNT 211 THERE ARE 211 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L9 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
- TI Fast dissolving orally consumable films containing an ion exchange resin as a taste masking agent
- AΒ Physiol. acceptable films, including edible films, are disclosed. films include a water soluble film-forming polymer, such as pullulan, and a taste masked pharmaceutically active agent, such as dextromethorphan. The taste masking agent is preferably a sulfonated polymer ion exchange resin comprising polystyrene cross-linked with divinylbenzene, such as Amberlite. Methods for producing the films are also disclosed. For example, an antitussive film was prepared in accordance with the following procedure: (A) uncoated dextromethorphan hydrobromide was dissolved with mixing in the water, while maintaining the temperature at  $75\,^{\circ}$ , Amberlite resin was then mixed into the water with heating at  $70-80^{\circ}$ , and heating was stopped, water lost to evaporation was replaced, and the potassium sorbate and sweeteners were then added to the composition with mixing to form Preparation A. (B) The film-forming ingredients (i.e., xanthan gum, locust bean gum, carrageenan and pullulan) were mixed in a sep. container to form Preparation B. (C) Preparation B was slowly added to Preparation A with rapid mixing,

followed by overnight mixing at a reduced rate to provide Preparation C. (D)
The menthol was dissolved with mixing in the alc. in a sep. container.
The Physcool was then dissolved with mixing therein. Monoammonium
glycyrrhizinate, Polysorbate 80, Atmos 300 and flavors were then added to
the mixture and mixed to enhanced uniformity to form Preparation D. (E)
Preparation

D, glycerin and mannitol were added to Preparation C with thorough mixing to provide Preparation E. Preparation E was poured on a mold and cast to form a film

of a desired thickness at room temperature. The film was dried under warm air and cut to a desired dimension (dictated by, e.g., dosage and mouthfeel) for taste testing. The active film had a pleasing appearance and taste.

- AN 2001:713109 HCAPLUS <<LOGINID::20081006>>
- DN 135:262242
- ${\tt TI}$  Fast dissolving orally consumable films containing an ion exchange resin as a taste masking agent
- IN Bess, William S.; Kulkarni, Neema; Ambike, Suhas H.; Ramsay, Michael Paul
- PA Warner-Lambert Company, USA
- SO PCT Int. Appl., 41 pp.
- CODEN: PIXXD2
- DT Patent
- LA English
- FAN.CNT 1

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WO 2001-US2192 W 20010123 <--
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RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS BECOME

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS BECOME
                      IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                       ALL CITATIONS AVAILABLE IN THE RE FORMAT
 L9
         ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
 ΤI
         Fast dissolving orally consumable films
 AΒ
         Physiol. acceptable films, including edible films, are disclosed. The
         films include a water soluble film-forming polymer such as pullulan. Edible
         films are disclosed that include pullulan and antimicrobially effective
         amts. of the essential oils thymol, Me salicylate, eucalyptol and menthol.
         The edible films are effective at killing the plaque-producing germs that
         cause dental plaque, gingivitis and bad breath. The film can also contain
         pharmaceutically active agents. Methods for producing the films are also
         disclosed.
         2000:227470 HCAPLUS <<LOGINID::20081006>>
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        132:255811
         Fast dissolving orally consumable films
 ΤI
         Leung, Sau-Hung Spence; Leone, Robert S.; Kumar, Lori Dee; Kulkarni,
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         Neema; Sorg, Albert F.
 PA
         Warner-Lambert Company, USA
 SO
         PCT Int. Appl., 54 pp.
         CODEN: PIXXD2
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         English
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- L9 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2008 ACS on STN
- TI Drug interaction prevention in combination dose units
- AB A combination therapy dose unit designed to prevent interaction between a plurality of active agents is prepared by charging particles of an active agent, charging particles of an inert particulate medium with a charge of opposite polarity, and allowing the charged particulate medium particles to adhere electrostatically to the charged particles of the active agent, thereby coating the active agent with inert particulate medium. Thereafter, other active agents can be treated in a similar manner and the electrostatically coated active agents can be combined, optionally together with other noncoated active agents, into a single combination

therapy dose unit such as a tablet. Thus, microparticles containing Bi subsalicylate, PVP (binder), lactose (filler), and an exploder were passed over a pos. electrode at 20,000-30,000 V and 50-120 mA to render the microparticle surface pos. charged. Micronized Mg stearate (inert particulate medium) was neg. charged by passing it over a neg. electrode in a similar manner, and was then used to form a microscopic coat around the pos. charged particles. Microparticles of tetracycline-HCl and of metronidazole were similarly pos. charged and coated with inert particulate medium, and all 3 coated active agents then blended, combined with binders, fillers, and disintegrants, and compressed into tablets containing Bi, tetracycline-HCl, and metronidazole in proportions of 100:200:200.

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AN 1996:273495 HCAPLUS <<LOGINID::20081006>>
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DN 124:298965

OREF 124:55219a,55222a

TI Drug interaction prevention in combination dose units

IN Moore, Trevor; Borody, Thomas Julius

PA Australia

SO PCT Int. Appl., 15 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

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